Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A pharmaceutical composition comprising Eudragit 4135F present in an amount of about 20 to 90% w/w; a lubricant present in an amount of 0 to about 30% w/w; a dissolution modifying excipient present in an amount of about 2.5 to about 70% w/w, and optionally a surfactant present in an amount of 0 to 10%, a plasticizer present in an amount of 0 to 10% w/w and/or a processing agent present in an amount of 0 to about 10% w/w ~~for moulded capsule, shell or linker components~~.

2. (Original) The composition according to Claim 1 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w.

3. (Original) The composition according to Claim 1 which comprises a surfactant which is present in an amount of less than 5% w/w.

4. (Original) The composition according to Claim 3 wherein the surfactant is sodium dodecyl sulphate or is a block copolymer of ethylene oxide and propylene oxide.

5. (Original) The composition according to Claim 4 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

6. (Original) The composition according to Claim 4 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide.

7. (Original) The composition according to Claim 1 wherein the lubricant is present in an amount of about 10 to 30% w/w.

8. (Previously presented) The composition according to Claim 1 wherein the lubricant is stearyl alcohol, glycerol monostearate (GMS), talc, magnesium stearate, silicon dioxide, amorphous silicic acid, or fumed silica; and combinations or mixtures thereof.

9. (Original) The composition according to Claim 8 wherein the lubricant is stearyl alcohol.

10. (Original) The composition according to Claim 9 wherein the stearyl alcohol is present from about 10 to about 15% w/w.

11. (Original) The composition according to Claim 1 wherein the dissolution modifying excipient is a swellable solid which is ethyl cellulose, cellulose acetate phthalate; hydroxypropyl cellulose, hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or other hydroxyalkylcellulose derivative; and combinations or mixtures thereof.

12. (Original) The composition according to Claim 11 wherein the dissolution modifying excipient is hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or hydroxypropyl cellulose.

13. (Currently amended) The composition according to Claim 12 wherein the swellable solid is present in an amount of about 10 to 50% w/w.

14. (Previously presented) The composition according to Claim 1 wherein the dissolution modifying excipient is xylitol, mannitol, lactose, pregelatinized starch sodium chloride, sodium starch glycollate, croscarmellose sodium, crospovidone (cross-linked polyvinyl pyrrolidone), copovidone , polyvinyl pyrrolidone, -and combinations or mixtures thereof.

15. (Previously presented) The composition according to Claim 14 wherein the dissolution modifying excipient is present in an amount of about 40 to 70% w/w.

16. (Previously presented) The composition according to Claim 11 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium

starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone.

17. (Original) The composition according to Claim 16 wherein the dissolution modifying excipient is hydroxypropylcellulose and lactose.

18. (Original) The composition according to Claim 1 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide, lecithin, sodium dioctyl sulfosuccinate, sodium dodecyl sulphate, Polyoxyl 40 hydrogenated castor oil, polyoxyethylene sorbitan fatty acid esters, the sorbitan fatty acid esters, polyethylene glycol, Vitamin E-TPGS® (d-alpha-tocopheryl polyethylene glycol 1000 succinate), sucrose fatty acid ester; and combinations and mixtures thereof.

19. (Previously presented) The composition according Claim 18 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone.

20. (Original) The composition according to Claim 1 wherein the plasticizer is triethyl citrate (TEC), tributyl citrate, acetyl triethyl citrate (ATEC), acetyl tributyl citrate (ATBC), dibutyl phthalate, dibutyl sebacate (DBS), diethyl phthalate, vinyl pyrrolidone glycol triacetate, polyethylene glycol, polyoxyethylene sorbitan monolaurate, propylene glycol, or castor oil; and combinations or mixtures thereof.

21. (Original) The composition according to Claim 1 wherein the processing agent is talc.

22. (Original) The composition according to Claim 21 wherein the processing agent is present in an amount of about 1 to about 5 % w/w.

23. (Original) The composition according to Claim 1 which further comprises an absorption enhancer.

24. (Original) The composition according to Claim 23 wherein the absorption enhancer is chitosan, lecithin, lectin, a sucrose fatty acid ester, Vitamin E-TPGS; and combinations or mixtures thereof.

25. (Original) The composition according to Claim 1 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w, the lubricant is stearyl alcohol, and the dissolution modifying excipient is hydroxypropylmethylcellulose, hydroxypropylcellulose, or a hydroxylalkyl cellulose derivative or salt thereof.

26. (Original) The composition according to Claim 25 wherein the dissolution modifying excipient also includes a disintegrant.

27. (Previously presented) The composition according to Claim 26 wherein the disintegrant is sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone, or a combination or mixture thereof.

28. (Original) The composition according to Claim 25 wherein the dissolution modifying excipient also includes a wicking agent.

29. (Original) The composition according to Claim 28 wherein the wicking agent is lactose.

30. (Original) The composition according to Claim 25 wherein the processing aid is talc.

31. (Previously presented) The pharmaceutical composition according to Claim 1 which is:

Example #	Formulation	% w/w
1.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Croscarmellose sodium	20.0
2.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	20.0
3.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Xylitol	10.0
4.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Croscarmellose sodium	10.0
	Xylitol	10.0
5.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Mannitol	10.0
	Sodium starch glycollate	10.0
6.	Eudragit 4135F	65.0
	Stearyl alcohol	5.0
	Mannitol	10.0
	Sodium starch glycollate	20.0

7.	Eudragit 4135F	80.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	10.0
	Lactose monohydrate	5.0
8.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	10.0
	Lactose monohydrate	10.0
9.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Lactose monohydrate	10.0
10.	Eudragit 4135F	75.0
	Stearyl alcohol	5.0
	Lactose monohydrate	20.0
11.	Eudragit 4135F	80.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	5.0
	Lactose monohydrate	10.0
12.	Eudragit 4135F	70.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	5.0
	Lactose monohydrate	20.0
13.	Eudragit 4135F	75.0
	Stearyl alcohol	10.0
	Mannitol	7.5
	Sodium starch glycollate	7.5
14.	Eudragit 4135F	80.0
	Stearyl alcohol	5.0
	Pregelatinized Starch	10.0
15.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0

	Pregelatinized Starch	15.0
16.	Eudragit 4135F	80.0
	Stearyl alcohol	5.0
	Pregelatinized Starch	10.0
	Lactose monohydrate	5.0
17.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Cross linked polyvinyl pyrrolidone	10.0
18.	Eudragit 4135F	80.0
	Stearyl alcohol	5.0
	Sodium starch glycollate	10.0
	Lactose monohydrate	5.0
19.	Eudragit 4135F	75.0
	Stearyl alcohol	10.0
	Sodium starch glycollate	10.0
	Lactose monohydrate	5.0
20.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Sodium chloride	5.0
	Lactose monohydrate	5.0
21.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Hydroxypropyl cellulose	5.0
	Lactose monohydrate	5.0
22.	Eudragit 4135F	85.0
	Stearyl alcohol	5.0
	Hydroxypropylmethyl cellulose	5.0
	Lactose monohydrate	5.0
23.	Eudragit 4135F	80.0
	Stearyl alcohol	10.0
	Hydroxypropylmethyl cellulose	5.0
	Lactose monohydrate	5.0

24.	Eudragit 4135F	80.0
	Stearyl alcohol	10.0
	Sodium starch glycollate	5.0
	Lactose monohydrate	5.0
25.	Eudragit 4135F	80.0
	Stearyl alcohol	10.0
	Hypromellose phthallate	5.0
	Lactose monohydrate	5.0
26.	Eudragit 4135F	80.0
	Stearyl alcohol	10.0
	Low substituted hydroxypropyl cellulose	5.0
	Lactose monohydrate	5.0
27.	Eudragit 4135F	90.0
	Stearyl alcohol	5.0
	Hydroxypropylmethyl cellulose	5.0
28.	Eudragit 4135F	90.0
	Stearyl alcohol	5.0
	Lactose monohydrate	5.0
29.	Eudragit 4135F	73.0
	Stearyl alcohol	12.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose monohydrate	5.0
30.	Eudragit 4135F	84.0
	Sodium dodecyl sulphate	1.0
	Croscarmellose sodium	15%

31.	Eudragit 4135F Sodium dodecyl sulphate Croscarmellose sodium Sodium starch glycollate	79.0 1.0 10% 10%
32.	Eudragit 4135F Croscarmellose sodium Sodium starch glycollate	80.0 10% 10%
33.	Eudragit 4135F Sodium dodecyl sulphate Croscarmellose sodium Sodium starch glycollate	69.0 1.0 15% 15%
34.	Eudragit 4135F Polyoxypropylene-polyoxyethylene block copolymer Sodium starch glycollate	79.0 1.0 20%
35.	Eudragit 4135F Polyoxypropylene-polyoxyethylene block copolymer Sodium starch glycollate	79.0 1.0 20%

32. (Previously presented) A pharmaceutical composition for molded capsule shells comprising:

Components	# (1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w	(7) w/w
Eudragit 4135F	45%	35%	25%	15%	75%	65%	55%
Stearyl Alcohol	10%	10%	10%	10%	10%	10%	10%
Lactose	5%	5%	5%	5%	5%	5%	5%
Hydroxypropyl Cellulose	40%	50%	60%	70%	10%	20%	30%
Total	100%	100%	100%	100%	100%	100%	100%

33. (Previously presented) A pharmaceutical composition for molded capsule shells comprising:

Components	# (1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w
Eudragit 4135F	63%	62.9%	62.75%	52%	42%	62%
Croscarmellose sodium	10%	10%	10%	15%	20%	5%
Sodium starch glycollate	10%	10%	10%	15%	20%	5%
Stearyl alcohol	12%	12%	12%	12%	12%	12%
Hydroxypropyl-methylcellu	5%	5%	5%	5%	5%	15%
Sodium Dodecyl Sulphate	0%	0.1%	0.25%	1%	1%	1%

34. (Previously presented) A pharmaceutical composition for molded capsule shells comprising:

Components	#(1) w/w	(2) w/w	(3) w/w	(4) w/w	(5) w/w	(6) w/w	(7) w/w
Eudragit 4135F	45%	35%	25%	15%	75%	65%	55%
Stearyl Alcohol	10%	10%	10%	10%	10%	10%	10%
Lactose	5%	5%	5%	5%	5%	5%	5%
Hydroxypropyl-cellulose	40%	50%	60%	70%	10%	20%	30%
Total	100%	100%	100%	100%	100%	100%	100%

35. (Previously presented) A pharmaceutical composition for molded capsule shells comprising:

Example #	Formulation	%w/w
1.	Eudragit 4135F	73.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Glyceryl monostearate	12.0
2.	Eudragit 4135F	53.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Hydroxypropylmethyl cellulose phthallate	20.0
	Stearyl alcohol	12.0
3.	Eudragit 4135F	20.0
	Hydroxypropylmethyl cellulose	10.0
	Hydroxypropylmethyl cellulose phthallate	20.0
	Stearyl alcohol	12.0
4.	Eudragit 4135F	68.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	5.0
	Stearyl alcohol	12.0
5.	Eudragit 4135F	72.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
6.	Eudragit 4135F	71.0
	Hydroxypropylmethyl cellulose	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	2.0

	Stearyl alcohol	12.0
7.	Eudragit 4135F	62.0
	Sodium starch glycollate	20.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
8.	Eudragit 4135F	62.0
	Sodium starch glycollate	20.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
9.	Eudragit 4135F	72.0
	Sodium starch glycollate	10.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
10.	Eudragit 4135F	62.0
	Croscarmellose sodium	20.0
	Lactose (regular)	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
11.	Eudragit 4135F	62.0
	Sodium starch glycollate	20.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
12.	Eudragit 4135F	62.0
	Hydroxypropylmethyl cellulose	20.0
	phthallate	5.0
	Hydroxypropylmethyl cellulose	1.0
	Sodium dodecyl sulphate	12.0
	Stearyl alcohol	

13.	Eudragit 4135F	62.5
	Sodium starch glycollate	20.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	0.5
	Stearyl alcohol	12.0
14.	Eudragit 4135F	62.0
	Croscarmellose sodium	10.0
	Sodium starch glycollate	10.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
15.	Eudragit 4135F	67.0
	Croscarmellose sodium	15.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
16.	Eudragit 4135F	72.0
	Croscarmellose sodium	10.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
17.	Eudragit 4135F	77.0
	Croscarmellose sodium	5.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
18.	Eudragit 4135F	52.0
	Croscarmellose sodium	15.0
	Sodium starch glycollate	15.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0

19.	Eudragit 4135F	42.0
	Croscarmellose sodium	20.0
	Sodium starch glycollate	20.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
20.	Eudragit 4135F	42.0
	Croscarmellose sodium	20.0
	Sodium starch glycollate	20.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
21.	Eudragit 4135F	62.0
	Croscarmellose sodium	5.0
	Sodium starch glycollate	5.0
	Hydroxypropylmethyl cellulose	15.0
	Sodium dodecyl sulphate	1.0
	Stearyl alcohol	12.0
22.	Eudragit 4135F	62.9
	Croscarmellose sodium	10.0
	Sodium starch glycollate	10.0
	Hydroxypropylmethyl cellulose	5.0
	Sodium dodecyl sulphate	0.1
	Stearyl alcohol	12.0

36. (Cancelled)

37. (Cancelled)

38. (Previously presented) The composition according to Claim 1 wherein the lubricant is stearyl alcohol present in an amount of 10 to 15% w/w, the surfactant is SDS or a block copolymer of ethylene oxide and propylene oxide present in an amount less than 5% w/w; a dissolution modifying excipient selected from HPC, HPMC, sodium starch glycollate,

croscarmellose sodium, copovidone, or lactose, and combinations or mixtures thereof, present in an amount of about 2.5 to about 70% w/w.

39. (Previously presented) A composition according to Claim 1 that is in the form of an injection molded capsule shell, linker or spacer.

40. (Previously presented) A composition according to Claim 1 that is in the form of a multicomponent injection molded capsule shell, linker or spacer.

41. (Previously presented) A composition according to Claim 1 that is in the form of a welded multicomponent injection molded capsule shell, linker or spacer.

42 to 70 (cancelled).

71. (Previously presented) The composition according to Claim 1 wherein the

	Dissolution Modifier	Lubricant	Surfactant
1.	Hydroxypropylmethylcellulose (5%w/w)	Stearyl alcoho (12%w/w)	None; or
2.	Hydroxypropylmethylcellulose (10%w/w), and HPMCphthalate (20%w/w)	Stearyl alcoho (12%w/w)	None; or
3.	Hydroxypropylmethylcellulose (10%), and Lactose (5%)	Stearyl alcoho (12%)	None; or
4.	Hydroxypropylmethylcellulose (	Stearyl alcohol (12%)	SDS (1%) or Sodium Starch Glycollate (20%) or Tween or a polyoxypropylene-polyoxyethylene block copolymer.

72. (Previously presented) The composition according to Claim 1 which is

Example #	Formulation	% w/w
1.	Eudragit 4135F	77.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	5.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
2.	Eudragit 4135F	68.0
	Croscarmellose sodium	15.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
3.	Eudragit 4135F	62.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	10.0
	Sodium Starch Glycollate	10.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
4.	Eudragit 4135F	63.0
	Croscarmellose sodium	10.0
	Sodium Starch Glycollate	10.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
5.	Eudragit 4135F	52.0
	Sodium Dodecyl Sulphate	1.0
	Croscarmellose sodium	15.0
	Sodium Starch Glycollate	15.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0

6.	Eudragit 4135F	62.0
	Polyoxypropylene-polyoxyethylene block copolymer	1.0
	Sodium Starch Glycollate	20.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
7.	Eudragit 4135F	62.0
	polyoxypropylene-polyoxyethylene block copolymer	1.0
	Sodium Starch Glycollate	20.0
	Stearyl Alcohol	12.0
	Hydroxypropylmethyl Cellulose	5.0
8.	Eudragit 4135F	62.0
	Stearyl Alcohol	12.0
	Croscarmellose sodium	5.0
	Sodium Starch Glycollate	5.0
	Hydroxypropylmethyl Cellulose	15.0
	Sodium Dodecyl Sulphate	1.0
9.	Eudragit 4135F	42.0
	Stearyl Alcohol	12.0
	Croscarmellose sodium	20.0
	Sodium Starch Glycollate	20.0
	Hydroxypropylmethyl Cellulose	5.0
	Sodium Dodecyl Sulphate	1.0
10.	Eudragit 4135F	47.0
	Stearyl Alcohol	12.0
	Sodium Starch Glycollate	10.0
	Hydroxypropylmethyl Cellulose	30.0
	Sodium Dodecyl Sulphate	1.0

73. (Previously presented) A pharmaceutical composition comprising Eudragit 4135F present in an amount of about 20 to 90% w/w; a lubricant present in an amount of 10 to about 30% w/w; a dissolution modifying excipient present in an amount of about 2.5 to about 70% w/w, and optionally a surfactant present in an amount of less than 5% w/w, a plasticizer present in an amount of 0 to 10% w/w and/or a processing agent present in an amount of 0 to about 10% w/w for moulded capsule, shell or linker components.

74. (Previously presented) The composition according to Claim 73 wherein the Eudragit 4135F is present in an amount of about 50 to 90% w/w.

75. (Previously presented) The composition according to Claim 73 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide, lecithin, sodium dioctyl sulfosuccinate, sodium dodecyl sulphate, Polyoxyl 40 hydrogenated castor oil, polyoxyethylene sorbitan fatty acid esters, the sorbitan fatty acid esters, polyethylene glycol, Vitamin E-TPGS® (d-alpha-tocopheryl polyethylene glycol 1000 succinate), sucrose fatty acid ester; and combinations and mixtures thereof.

76. (Previously presented) The composition according to Claim 75 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

77. (Previously presented) The composition according to Claim 75 wherein the surfactant is sodium dodecyl sulphate or is a block copolymer of ethylene oxide and propylene oxide.

78. (Previously presented) The composition according to Claim 77 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

79. (Previously presented) The composition according to Claim 73 wherein the surfactant is a block copolymer of ethylene oxide and propylene oxide.

80. (Previously presented) The composition according to Claim 79 wherein the surfactant is sodium dodecyl sulphate is present in an amount of less than 2% w/w.

81. (Previously presented) The composition according to Claim 73 wherein the lubricant is stearyl alcohol, glycerol monostearate (GMS), talc, magnesium stearate, silicon dioxide, amorphous silicic acid, or fumed silica; and combinations or mixtures thereof.

82. (Previously presented) The composition according to Claim 81 wherein the lubricant is stearyl alcohol.

83. (Previously presented) The composition according to Claim 82 wherein the stearyl alcohol is present from about 10 to about 15% w/w.

84. (Previously presented) The composition according to Claim 73 wherein the dissolution modifying excipient is a swellable solid which is ethyl cellulose, cellulose acetate phthalate; hydroxypropyl cellulose, hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or other hydroxyalkylcellulose derivative; and combinations or mixtures thereof.

85. (Previously presented) The composition according to Claim 84 wherein the swellable solid is present in an amount of about 10 to 50% w/w.

86. (Previously presented) The composition according to Claim 84 wherein the dissolution modifying excipient is hydroxypropylmethyl cellulose, hydroxypropylmethyl cellulose phthalate, or hydroxypropyl cellulose.

87. (Previously presented) The composition according to Claim 86 wherein the swellable solid is present in an amount of 10 to 50% w/w.

88. (Previously presented) The composition according to Claim 73 wherein the dissolution modifying excipient is xylitol, mannitol, lactose, pregelatinized starch, sodium chloride, sodium starch glycollate, croscarmellose sodium, crospovidone (cross-linked polyvinyl pyrrolidone), copovidone, polyvinyl pyrrolidone; and combinations or mixtures thereof.

89. (Previously presented) The composition according to Claim 88 wherein the dissolution modifying excipient is present in an amount of about 40 to 70% w/w.

90. (Previously presented) The composition according to Claim 89 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, crospovidone (cross-linked polyvinyl pyrrolidone), or polyvinyl pyrrolidone.

91. (Previously presented) The composition according to Claim 90 wherein the dissolution modifying excipient is hydroxypropylcellulose and lactose.

92. (Previously presented) The composition according Claim 73 wherein the dissolution modifying excipient is a combination of a swellable solid and lactose, sodium starch glycollate, croscarmellose sodium, copovidone, or crospovidone (cross-linked polyvinyl pyrrolidone).

93. (Previously presented) The composition according to Claim 73 wherein the processing agent is talc.

94. (Previously presented) The composition according to Claim 93 wherein the processing agent is present in an amount of about 1 to about 5 % w/w.

95. (Previously presented) The composition according to Claim 91 wherein the processing agent is talc and is present in an amount of about 1 to about 5 % w/w.

96. (Previously presented) The composition according to Claim 73 which further comprises an absorption enhancer.

97. (Previously presented) The composition according to Claim 96 wherein the absorption enhancer is chitosan, lecithin, lectin, a sucrose fatty acid ester, Vitamin E-TPGS; and combinations or mixtures thereof.

98. (Currently amended) A pharmaceutical composition comprising Eudragit 4135F is present in an amount of about 50 to about 90% w/w, the lubricant is stearyl alcohol, present in an amount of about 10 to about 15% w/w, and ~~the~~ a dissolution modifying excipient selected from ~~is~~ hydroxypropylmethylcellulose, hydroxypropylcellulose, or a hydroxylalkyl cellulose derivative or salt thereof ~~, for moulded capsule, shell, or linker components.~~

99. (Previously presented) The composition according to Claim 98 wherein the dissolution modifying excipient also includes a disintegrant.

100. (Previously presented) The composition according to Claim 99 wherein the disintegrant is sodium starch glycollate, croscarmellose sodium, copovidone, or crospovidone (cross-linked polyvinyl pyrrolidone), or a combination or mixture thereof.

101. (Previously presented) The composition according to Claim 100 wherein the dissolution modifying excipient also includes a wicking agent.

102. (Previously presented) The composition according to Claim 101 wherein the wicking agent is lactose.

103. (Previously presented) The composition according to Claim 101 which further comprises a processing aid which is talc.

104. (Previously presented) The composition according to Claim 98 wherein the dissolution modifying excipient also includes a wicking agent.

105. (Previously presented) The composition according to Claim 104 wherein the wicking agent is lactose.

106. (Previously presented) The composition according to Claim 98 wherein the HPC is present in an amount of 10 to about 70% w/w.

107. (Previously presented) The composition according to Claim 106 wherein the HPC is present in an amount of 40 to about 70% w/w.

108. (Previously presented) The composition according to Claim 107 further comprises a wicking agent which is lactose.

109. (Previously presented) The composition according to Claim 107 wherein the lactose is present in an amount of about 0 to 10% w/w.

110. (Previously presented) The composition according to Claim 107 wherein the lactose is present in an amount of about 5% w/w.

111. (Currently amended) The composition according to Claim 99 wherein the disintegrant ~~distintegrant~~ is present in an amount of about 10 to about 40% w/w.